

WHAT IS CLAIMED IS:

1           1. An isolated nucleic acid encoding a G-protein coupled receptor  
2 polypeptide, the nucleic acid encoding a polypeptide comprising greater than 70% amino  
3 acid identity to an amino acid sequence of SEQ ID NO:6, SEQ ID NO:4, SEQ ID NO:8,  
4 SEQ ID NO:10, SEQ ID NO:12, or SEQ ID NO:16.

*Sub B1*  
1           2. An isolated nucleic acid of claim 1, wherein the nucleic acid  
2 encodes a polypeptide comprising greater than 80% amino acid identity to an amino acid  
3 sequence of SEQ ID NO:6, SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:10, SEQ ID  
4 NO:12, or SEQ ID NO:16.

1           3. An isolated nucleic acid of claim 1, wherein the nucleic acid  
2 encodes a polypeptide comprising greater than 90% amino acid identity to an amino acid  
3 sequence of SEQ ID NO:6, SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:10, SEQ ID  
4 NO:12, or SEQ ID NO:16.

1           4. The isolated nucleic acid of claim 1, wherein the nucleic acid  
2 encodes a polypeptide that specifically binds to polyclonal antibodies generated against  
3 an amino acid sequence of SEQ ID NO:6, SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:10,  
4 SEQ ID NO:12, or SEQ ID NO:16.

1           5. The isolated nucleic acid of claim 1, wherein the nucleic acid  
2 encodes a polypeptide that has G-protein coupled receptor activity.

*Sub B2*  
1           6. The isolated nucleic acid of claim 1, wherein the nucleic acid  
2 encodes a polypeptide comprising an amino acid sequence of SEQ ID NO:6, SEQ ID  
3 NO:4, SEQ ID NO:8, SEQ ID NO:10, SEQ ID NO:12, or SEQ ID NO:16.

1           7. The isolated nucleic acid of claim 1, wherein the nucleic acid  
2 comprises the nucleotide sequence of SEQ ID NO:5, SEQ ID NO:3, SEQ ID NO:7, SEQ  
3 ID NO:9, SEQ ID NO:11, or SEQ ID NO:15.

1           8. The isolated nucleic acid of claim 1, wherein the nucleic acid is  
2 amplified by primers that specifically hybridize under stringent hybridization conditions  
3 to a nucleic acid having a nucleotide sequence of SEQ ID NO:5, SEQ ID NO:3, SEQ ID  
4 NO:7, SEQ ID NO:9, SEQ ID NO:11, or SEQ ID NO:15.

08860

1           9. An isolated nucleic acid encoding a G-protein coupled receptor  
2 polypeptide, wherein the nucleic acid specifically hybridizes under stringent hybridization  
3 conditions to a nucleic acid having a nucleotide sequence of SEQ ID NO:5, SEQ ID  
4 NO:3, SEQ ID NO:7, SEQ ID NO:9, SEQ ID NO:11, or SEQ ID NO:15.

1           10. An isolated nucleic acid encoding a G-protein coupled receptor  
2 polypeptide, the polypeptide encoded by the nucleic acid comprising greater than about  
3 70% amino acid identity to a polypeptide having an amino acid sequence of SEQ ID  
4 NO:6, SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:10, SEQ ID NO:12, or SEQ ID NO:16,  
5 wherein the nucleic acid selectively hybridizes under moderately stringent hybridization  
6 conditions to a nucleotide sequence of SEQ ID NO:5, SEQ ID NO:3, SEQ ID NO:7, SEQ  
7 ID NO:9, SEQ ID NO:11, or SEQ ID NO:15.

1           11. An isolated nucleic acid encoding a G-protein coupled receptor  
2 polypeptide, wherein the nucleic acid encodes a polypeptide comprising at least 25  
3 contiguous amino acids of the amino acid sequence of SEQ ID NO:6, SEQ ID NO:4,  
4 SEQ ID NO:8, SEQ ID NO:10, SEQ ID NO:12, or SEQ ID NO:16.

1           12. The isolated nucleic acid of claim 11, wherein the nucleic acid  
2 encodes a polypeptide that comprises at least 50 contiguous amino acids of the amino  
3 acid sequence of SEQ ID NO:6, SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:10, SEQ ID  
4 NO:12, or SEQ ID NO:16.

1           Sub 33 13. An isolated nucleic acid encoding a G-protein coupled receptor  
2 polypeptide, wherein the nucleic acid encodes a polypeptide comprising greater than 90%  
3 amino acid identity to an amino acid sequence of SEQ ID NO:2 or SEQ ID NO:14.

1           14. The isolated nucleic acid of claim 13, wherein the nucleic acid  
2 encodes a polypeptide that specifically binds to polyclonal antibodies generated against  
3 an amino acid sequence of SEQ ID NO:2 or SEQ ID NO:14.

1           15. The isolated nucleic acid of claim 13, wherein the nucleic acid  
2 encodes a polypeptide that has G-protein coupled receptor activity.

1           16. The isolated nucleic acid of claim 13, wherein the nucleic acid  
2 encodes a polypeptide comprising an amino acid sequence of SEQ ID NO:2 or SEQ ID  
3 NO:14.

1           17. The isolated nucleic acid of claim 13, wherein the nucleic acid  
2 comprises the nucleotide sequence of SEQ ID NO:1 or SEQ ID NO:13.

1           18. An isolated nucleic acid encoding a G-protein coupled receptor  
2 polypeptide, the polypeptide encoded by the nucleic acid comprising greater than about  
3 90% amino acid identity to a polypeptide having an amino acid sequence of SEQ ID  
4 NO:2 or SEQ ID NO:14, wherein the nucleic acid selectively hybridizes under  
5 moderately stringent hybridization conditions to a nucleotide sequence of SEQ ID NO:1  
6 or SEQ ID NO:13.

1           19. An isolated G-protein coupled receptor polypeptide, the  
2 polypeptide comprising greater than about 70% amino acid sequence identity to an amino  
3 acid sequence of SEQ ID NO:6, SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:10, SEQ ID  
4 NO:12, or SEQ ID NO:16.

1           20. The isolated polypeptide of claim 19, wherein the polypeptide  
2 comprises greater than 80% amino acid sequence identity to an amino acid sequence of  
3 SEQ ID NO:6, SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:10, SEQ ID NO:12, or SEQ ID  
4 NO:16.

1           21. The isolated polypeptide of claim 19, wherein the polypeptide  
2 comprises greater than 90% amino acid sequence identity to an amino acid sequence of  
3 SEQ ID NO:6, SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:10, SEQ ID NO:12, or SEQ ID  
4 NO:16.

1           22. The isolated polypeptide of claim 19, wherein the polypeptide  
2 specifically binds to polyclonal antibodies generated against SEQ ID NO:6, SEQ ID  
3 NO:4, SEQ ID NO:8, SEQ ID NO:10, SEQ ID NO:12, or SEQ ID NO:16.

1           23. The isolated polypeptide of claim 19, wherein the polypeptide has  
2 G-protein coupled receptor activity.

1           24. The isolated polypeptide of claim 19, wherein the polypeptide has  
2 the amino acid sequence of SEQ ID NO:6, SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:10,  
3 SEQ ID NO:12, or SEQ ID NO:16.

1           25. An isolated G-protein coupled receptor polypeptide, the  
2 polypeptide comprising greater than about 90% amino acid sequence identity to an amino  
3 acid sequence of SEQ ID NO:2 or SEQ ID NO:14.

1           26. The isolated polypeptide of claim 25, wherein the polypeptide  
2 specifically binds to polyclonal antibodies generated against SEQ ID NO:2 or SEQ ID  
3 NO:14.

1           27. The isolated polypeptide of claim 25, wherein the polypeptide has  
2 G-protein coupled receptor activity.

1           28. The isolated polypeptide of claim 25, wherein the polypeptide has  
2 an amino acid sequence of SEQ ID NO:2 or SEQ ID NO:14.

1           29. An antibody that selectively binds to the polypeptide of claim 19,  
2 or 25.

1           30. An expression vector comprising the nucleic acid of claim 1, 11, or  
2 13.

1           31. A host cell transfected with the vector of claim 30.

1           32. A method for identifying a compound that modulates signal  
2 transduction, the method comprising:

3           (i) contacting the compound with a polypeptide comprising greater than  
4 70% amino acid sequence identity to the amino acid sequence of SEQ ID NO:6, SEQ ID  
5 NO:4, SEQ ID NO:8, SEQ ID NO:10, SEQ ID NO:12, or SEQ ID NO:16; and  
6           (ii) determining the functional effect of the compound upon the  
7 polypeptide.

1           33. The method of claim 32, wherein the polypeptide has G-protein  
2 coupled receptor activity.

1                   34.     The method of claim 32, wherein the polypeptide is linked to a  
2 solid phase.

1                   35.     The method of claim 34, wherein the polypeptide is covalently  
2 linked to a solid phase.

1                   36.     The method of claim 32, wherein the functional effect is  
2 determined by measuring changes in intracellular cAMP, IP<sub>3</sub>, or Ca<sup>2+</sup>.

1                   37.     The method of claim 32, wherein the functional effect is a chemical  
2 effect.

1                   38.     The method of claim 32, wherein the functional effect is a physical  
2 effect.

1                   39.     The method of claim 32, wherein the functional effect is  
2 determined by measuring binding of the compound to the polypeptide.

1                   40.     The method of claim 32, wherein the polypeptide is recombinant.

1                   41.     The method of claim 32, wherein the polypeptide comprises the  
2 amino acid sequence of SEQ ID NO:6, SEQ ID NO:4, SEQ ID NO:8, SEQ ID NO:10,  
3 SEQ ID NO:12, or SEQ ID NO:16.

1                   42.     The method of claim 32, wherein the polypeptide is expressed in a  
2 cell or cell membrane.

1                   43.     The method of claim 42, wherein the cell is a eukaryotic cell.

1                   44.     The method of claim 43, wherein the cell is an adipocyte.

1                   45.     The method of claim 43, wherein the cell is a spleen cell.

1                   46.     The method of claim 43, wherein the cell is a colon cell.

1                   47.     The method of claim 43, wherein the cell is a neuron.

1                   48.     A method for identifying a compound that modulates signal  
2 transduction, the method comprising the steps of:

PCT/US2007/034468

3 (i) contacting the compound with a polypeptide comprising greater than  
4 90% amino acid sequence identity to the amino acid sequence of SEQ ID NO:2 or SEQ  
5 ID NO:14; and

(ii) determining the functional effect of the compound upon the polypeptide.

1                          49.        The method of claim 48, wherein the polypeptide has G-protein  
2 coupled receptor activity.

1 50. The method of claim 48, wherein the polypeptide is linked to a  
2 solid phase.

51. The method of claim 48, wherein the functional effect is  
determined by measuring changes in intracellular cAMP, IP<sub>3</sub>, or Ca<sup>2+</sup>.

1                           52. The method of claim 48, wherein the functional effect is a chemical  
2                           effect.

1                           53.     The method of claim 48, wherein the functional effect is a physical  
2     effect.

1                   54. The method of claim 48, wherein the functional effect is  
2 determined by measuring binding of the compound to the polypeptide.

1 55. The method of claim 48, wherein the polypeptide is recombinant.

1                   56. The method of claim 48, wherein the polypeptide comprises the  
2 amino acid sequence of SEQ ID NO:2 or SEQ ID NO:14.

1                   57.     The method of claim 48, wherein the polypeptide is expressed in a  
2     cell or cell membrane.

1 58. The method of claim 57, wherein the cell is a eukaryotic cell.

1 59. The method of claim 58, wherein the cell is a kidney cell.

1                   60. A method of treating kidney disease, the method comprising the  
2 step of administering to a patient a therapeutically effective amount of a compound  
3 identified using the method of claim 48.

1               61. A method of treating cerebral cavernous malformations, the  
2 method comprising the step of administering to a patient a therapeutically effective  
3 amount of a compound identified using the method of claim 48.

1               62. A method of treating hyperlipidemia, the method comprising the  
2 step of administering to a patient a therapeutically effective amount of a compound  
3 identified using the method of claim 32.

1               63. A method of treating obesity, the method comprising the step of  
2 administering to a patient a therapeutically effective amount of a compound identified  
3 using the method of claim 32.

1               64. A method of treating dyslexia, the method comprising the step of  
2 administering to a patient a therapeutically effective amount of a compound identified  
3 using the method of claim 32.

1               65. A method of treating cardiac myxoma, the method comprising the  
2 step of administering to a patient a therapeutically effective amount of a compound  
3 identified using the method of claim 32.

1               66. A method of detecting the presence of an TGR-GPCR or a EDG-  
2 GPCR nucleic acid or polypeptide in human tissue, the method comprising the steps of:  
3                     (i) isolating a biological sample;  
4                     (ii) contacting the biological sample with a TGR-GPCR-specific  
5 reagent or a EDG-GPCR-specific reagent that selectively associates with an TRG-GPCR  
6 nucleic acid or polypeptide or a EDG-GPCR nucleic acid or polypeptide; and,  
7                     (iii) detecting the level of TGR-GPCR-specific reagent or EDG-  
8 GPCR-specific reagent that selectively associates with the sample.

1               67. The method of claim 66, wherein the TGR-GPCR-specific reagent  
2 or EDG-GPCR-specific reagent is selected from the group consisting of: antibodies,  
3 oligonucleotide primers, and nucleic acid probes.